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(54) Title: DETERMINING A PARAMETER OF A COMPONENT IN A COMPOSITION

#### (57) Abstract

A method of determining a parameter selected from the electrical conductivity and the volume fraction of a component in a composition comprising a plurality of components is provided. The method comprises measuring the electrical conductivity of the composition, and selecting a relationship between the conductivity of the composition and a plurality of composition parameters including, for each component, physical parameters representing the conductivity and the volume fraction of the component, said relationship being such that the components are substantially equally represented in said relationship by means of said physical parameters. The selected parameter of the component in the composition is determined from said relationship and the measured conductivity of the composition.

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# DETERMINING A PARAMETER OF A COMPONENT IN A COMPOSITION

The present invention relates to a method of determining a parameter selected from the electrical conductivity and the volume fraction of a component in a composition comprising a plurality of components. The invention is of particular interest for determining the volume fraction of a component of an earth formation, for example to determine the hydrocarbon-content of a hydrocarbon bearing earth formation. Various known methods of determining such content have been applied until now, which known methods are generally based on empirical models.

One such known method is described in "Electrical conductivities in oil-bearing shaly sands", Waxman M.H. and Smits L.J.M., SPE paper 1863-A presented at 42nd Ann. Fall Meeting, Houston, October 1-4, 1967.

This publication discloses a method of determining a parameter selected from the electrical conductivity and the volume fraction of a component in a composition comprising a plurality of components, wherein the electrical conductivity of the composition is measured, and a relationship between the conductivity of the composition and the conductivity of a component is selected.

This known method uses the following relationship which is generally referred to as the Waxman-Smits model:

$$C_O = C_W / F^* + BQ_V / F^*$$

where  $C_0$  = conductivity of fully brine saturated rock  $C_W = \text{conductivity of brine present in the formation}$ 

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- $F^* = a$  formation factor
- B = equivalent conductance of sodium clay-exchange cations as a function of  $C_{\mathbf{w}}$

 $Q_{\rm V}$  = cation exchange capacity per unit pore volume. The results achieved with this known method are not always sufficiently accurate, probably because of the empirical nature of the Waxman-Smits model which provides a relationship between the earth conductivity and the various other parameters.

It is an object of the invention to provide a more accurate method of determining a parameter selected from the electrical conductivity and the volume fraction of a component in a composition comprising a plurality of components.

The method according to the invention thereto comprises:

- measuring the electrical conductivity of the composition;
- selecting a relationship between the electrical conductivity of the composition and a plurality of composition parameters including, for each component, physical parameters representing the electrical conductivity and the volume fraction of the component, the components being substantially equally represented in said relationship by means of said physical parameters; and
- determining said selected parameter from said relationship and the measured conductivity of the composition.

It is to be understood that by the electrical conductivity is meant the electrical conductivity itself or any quantity derived therefrom, such as the electrical resistivity. Furthermore, the feature that the components are substantially equally represented in the relationship implies that each component is represented in the

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relationship in substantially the same way as any other of the components.

With the method according to the invention it is achieved that results of increased accuracy are provided. The selected relationship takes accurately account of the individual contributions of the components to the conductivity of the composition. The relationship applied in the method according to the invention is symmetrical in the components, i.e. no component is favoured over another component. Moreover, it was found that the method according to the invention provides the desired accuracy for any percolation threshold of the components. In this respect it is to be understood that the amount of percolation of a component refers to the degree of continuity of the component in the composition. For example, vanishing percolation of a component implies that the component is fully dispersed in the composition, and full percolation of a component implies that the component is continuous throughout the composition.

Advantageously said plurality of composition parameters includes at least one fitting parameter, and wherein each fitting parameter is determined by applying said relationship to a data set obtained by measuring the electrical conductivity of at least one sample representative for said composition for various magnitudes of at least one of said parameters.

Preferably said plurality of parameters includes an auxiliary parameter depending on the geometrical configuration of the components in the composition.

Accurate geometrical representation by the auxiliary component is achieved if said auxiliary parameter is selected so as to be a function of a plurality of variables, each variable depending on the conductivity of one of said components and a mixing coefficient, whereby

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the mixing coefficients depend on the geometrical configuration of the components in the composition.

Advantageously the step of determining each fitting parameter by applying said relationship to the data set of the component is carried out through an iterative process. Suitably the iterative process includes repeatedly applying said relationship in a minimisation scheme. The minimisation scheme is preferably applied to an incoherence between the measured electrical conductivities of said components and the electrical conductivities of the components as determined through said relationship.

The invention will be described hereinafter in more detail and by way of the following example and comparative example.

#### Example

Consider an isotropic system with essential spherical inclusions in the form of an earth formation which essentially consists of four components: non-conducting porous rock matrix, non-conducting hydrocarbon fluid, conducting clay, and conducting brine. The conductivity of the formation depends on the fractional brine saturation of the pore space, and the hydrocarbon fluid component is grouped with the rock matrix, both being non-conducting. Thus, the hydrocarbon component and the rock matrix component only enter the equations with the sum of their volume fractions. The effective conductivity  $\sigma_{\rm eff}$  of this earth formation is evaluated through the expression

$$(\sigma_{\text{eff}} - \sigma_0) \cdot (L\sigma_{\text{eff}} + (1-L)\sigma_0)^{-1} = \sum \phi_k (\sigma_k - \sigma_0) \cdot (L\sigma_k + (1-L)\sigma_0)^{-1}$$

wherein  $\sigma_0$  represents the auxiliary parameter in the form of a conductivity tensor

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 $k=1\ldots N$ , N being the number of components  $\sigma_{eff}$  represents the conductivity tensor of the sample  $\sigma_k$  represents the conductivity tensor of component k  $\phi_k$  represents the volume fraction of component k L represents the depolarisation tensor (shape tensor)

Preferably the depolarisation tensor is positive and has unit trace. In an attractive embodiment the depolarisation tensor equals 1/3 times the unit tensor.

The term  $\sigma_0$  denotes an auxiliary parameter which can be thought of as being an additional host medium into which components are added until the host medium has been completely replaced by the components so that no volume fraction is associated with the host medium. The existence of the host medium enables the model to be symmetrical in all its constituents: none of the components rock, clay or brine in the model is favoured over any of the other components. The dependence of  $\sigma_0$  on various parameters, yet to be determined, governs the percolation behaviour of the model. Setting  $\sigma_0 = \sigma_{brine}$ leads to the known Average T-matrix Approximation, also referred to as the generalised Clausius-Mossotti equation. This model has a clear asymmetry between the brine component and the other components since only the brine component will percolate, irrespective of its volume fraction. Selecting a self-consistent host medium conductivity,  $\sigma_0 = \sigma_{eff}$ , leads to the known Coherent Potential Approximation, also referred to as the generalised Bruggeman equation. This model is symmetrical in all components but has the drawback of requiring unrealistically high percolation thresholds for each component.

In a suitable embodiment, the auxiliary parameter  $\sigma_{\mathbf{0}}$  is selected as follows:

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$$\sigma_0 = \sum h_k \sigma_k$$
; for  $k = 1, 2, 3$ 

wherein  $h_k$  represents the mixing coefficient tensor pertaining to component k, which tensor contains mixing coefficients representing geometrical information on the spatial distribution of the components in the formation. These coefficients determine the connectivity, i.e. the amount of percolation of the individual components. The coefficients are non-negative and fulfil the normalisation condition:

$$\sum h_k = 1$$
; for  $k = 1, 2, 3$ 

The normalisation relation ensures that the resulting effective conductivity  $\sigma_{\text{eff}}$  satisfies the Hashin - Shtrikman bounds, which are well known to those skilled in the art.

Furthermore, a component with a vanishingly small volume fraction can not percolate, hence the corresponding connectivity parameter should vanish:

$$\lim h_k = 0$$
; for  $\phi_k \Rightarrow 0$ 

Suitably the mixing coefficient tensor is selected to be

$$h_k = \lambda_k \phi_k V_k \quad (\sum \lambda_n \phi_n V_n)^{-1}$$

wherein  $k, n = 1 \dots N$ , N being the number of components in said plurality of components

 $\lambda_{\boldsymbol{k}}$  represents the percolation rate tensor pertaining to component  $\boldsymbol{k}$ 

 $\varphi_k$  represents the volume fraction of component k  $\nu \quad \text{represents the percolation exponent pertaining to}$  component k

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Suitably at least one of  $h_k,\ \lambda_k$  and  $\nu$  forms a fitting parameter.

A data set on 27 shaly-sand core samples has been used to test the invention, which data set is described in the above indicated SPE paper. This publication provides Co - Cw curves on the core samples ranging from almost clean sand  $(Q_v = 0.017 \text{ eq/1})$  to extremely shaly sand  $(Q_V = 1.47 \text{ eq/l})$ . The samples contained Kaolinite, Montmorillonite and Illite, either in combination or separately in each sample. The characteristic petrophysical data of each sample are listed in the appended Table, in which  $\phi$  denotes the porosity of the sample,  $\kappa$  denotes the permeability of the sample, and  $Q_{\tau r}$ denotes the cation exchange capacity per unit pore volume of the sample. The conductivity of each sample in fully brine saturated condition was measured for eight to ten salinities of the brine. Furthermore, concentration membrane potential measurements were made of the samples.

The parameters in this model were selected as follows:

1) Brine;

The volume fraction of brine,  $\phi_D$ , is determined by the porosity, the amount of clay-bound water, and the water saturation  $S_W$ . The brine conductivity  $\sigma_D$  (= $C_W$ ) is determined by the brine salinity and the brine temperature. The two percolation parameters,  $\lambda_D$  and  $\nu$ , are free parameters.

2) Rock / Hydrocarbon;

The volume of hydrocarbons,  $\phi_{hc}$ , is determined by the total porosity, the amount of clay-bound water, and the hydrocarbon saturation 1-S<sub>W</sub>, while the volume of the rock matrix,  $\phi_r$ , is calculated using the sum rule and the volume fractions. Both the rock and the hydrocarbon have vanishing conductivity. The percolation parameters  $\lambda_r$  and  $\lambda_{hc}$  of both components was set at value 1. The mixing

coefficient pertaining to rock/hydrocarbon  $h_{r/hc}$  follows from the condition  $\sum h_k = 1$ .

3) Clay;

The volume of clay  $\phi_C$  and the clay conductivity  $\sigma_C$  are free fitting parameters. The percolation rate  $\lambda_C$  was set at a value 0, which is a suitable choice for non-laminated clays. It furthermore appeared that an additional free parameter did not give a significant improvement of the model fit to the data set.

The  $C_O - C_W$  measurements were made for an extreme salinity range, namely a brine salinity between 1 - 300 g/l. For a given sample the brine volume fraction varied only slightly over the whole salinity range. In view thereof the percolation parameter  $\nu$  was set equal to unity in the test, thereby reducing the percolation

free parameters to three. For each sample, a fit to the  $C_{\rm O}$  -  $C_{\rm W}$  curve was made by minimising the relative incoherence defined as:

parameter hb to a constant, and reducing the number of

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$$\sum \Delta^{2} C_{O} - C_{W} = \sum_{\text{salinities}} \begin{pmatrix} C_{O}, \text{calc}^{-C}_{O}, \text{meas} \\ ---- \\ C_{O}, \text{meas} \end{pmatrix}$$

wherein

Co,calc = the calculated conductivity of the fully brine
saturated rock samples;

Co,meas = the measured conductivity of the fully brine
saturated rock samples;

 $\Sigma$  = summation over the salinities.

The results for the three fitting parameters  $\phi_C,\ \sigma_C$  and  $h_D,$  and the relative incoherence are given in the appended Table.

Furthermore, the Table gives the incoherence between the membrane potential  $(\psi_{\text{Calc}})$  determined by the method of the invention and the measured membrane potential  $(\psi_{\text{meas}})$ :

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$$\sum \Delta^{2}_{MP} = \sum_{\text{salinities}} \begin{pmatrix} \psi_{\text{calc}} - \psi_{\text{meas}} \\ ----- \\ \psi_{\text{meas}} \end{pmatrix}$$
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The membrane potential is a particularly interesting quantity for being a direct, non-destructive, measure of the clay contribution to the overall conductivity, which has not been used to determine the fitting parameters.

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To illustrate the invention more specifically, reference is made to the following comparative example.

Comparative example

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As stated above Ref. 1 discloses, apart from the data set on the 27 core samples, furthermore an empirical model which is generally referred to as the Waxman-Smits model. To compare the method according to the invention with the Waxman-Smits model, the relative incoherence between the measured conductivities and the conductivities found from the Waxman-Smits model, and the relative incoherence between the measured concentration membrane potentials and the concentration membrane potentials found from the Waxman-Smits model, were determined. These relative incoherences for all 27 samples are listed in the Table. In applying the Waxman-Smits model, use has been made of the well known expression:

 $C_O = C_W / F^* + BQ_V / F^*$ with  $F^* = \phi^{-m}$ 

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where m is a free parameter (also referred to as the cementation exponent),  $Q_{\rm V}$  is determined from sample measurements, just as the porosity  $\phi$ , and the standard B-chart has been used to calculate the salinity and temperature effects on the conductivity measurements.

From a comparison between the incoherence values found by using the method according to the invention, and the incoherence values found by using the Waxman-Smits model, it is clear that the method according to the invention provides improved results. Especially the extremely low incoherence values for the concentration membrane potential, which values are moreover fairly constant over the entire  $Q_V$  range, indicates that the method according to the invention provides results of increased accuracy.

The method according to the invention can suitably be applied to determine the volume fraction of brine or hydrocarbon in an earth formation, whereby a well-log representing the electrical conductivity of the formation is provided. Such application can, for example, be carried out in the following manner. The well-log of the electrical conductivity of the earth formation is made using a logging tool lowered in a borehole formed in the earth formation. For an isotropic formation with components brine (subscript B), clay (subscript C), and non-conducting rock + hydrocarbon (subscript R/HC) the rock and the hydrocarbon are grouped together because of their vanishing conductivities. The selected relationship then is:

$$\sigma_{\text{eff}} - \sigma_{\text{o}} \qquad \qquad 3 \qquad \sigma_{\text{k}} - \sigma_{\text{o}}$$

$$----- = \sum \phi_{\text{k}}. \quad ----$$

$$\sigma_{\text{eff}} + 2 \sigma_{\text{o}} \qquad \qquad k=1 \qquad \sigma_{\text{k}} + 2\sigma_{\text{o}}$$

30 wherein

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$$\sigma_0 = \sum h_k \sigma_k$$
 $h_k = \lambda_k \phi_k^{\nu}_k (\sum \lambda_n \phi_n^{\nu})^{-1}$ 

in which

 $\sigma_0$  represents the auxiliary parameter

k, n = 1 ... N, N being the number components

 $\sigma_{\text{eff}}$  represents the conductivity of the earth formation

 $\sigma_k$  represents the conductivity of component k

 $\mathbf{t}_k$  represents the volume fraction of component k

h<sub>k</sub> represents the mixing coefficient pertaining to component k;

 $\boldsymbol{\lambda}_k$  represents the percolation coefficient pertaining to component k

 $v_{k,n}$  represents the percolation exponent pertaining to component k, n

Each component k has four parameters:  $\phi_k$ ,  $\sigma_k$ ,  $\lambda_k$ , and  $\nu_{k},$  of which  $\phi_{B},$   $\sigma_{B}$  and  $\sigma_{R/HC}$  are directly measured. Furthermore,  $\lambda_C = 0$  for dispersed clay. From the sum 15 rules  $h_{R/HC}$  and  $\phi_{R/HC}$  follow. Parameters which are yet to be determined are  $\sigma_C,~\lambda_B,~\nu_B$  and  $\varphi_C.$  These parameters are determined through forward modelling on experimental data.  $\sigma_C$ ,  $\lambda_B$  and  $\nu_B$  are invariable over the geological formation , while  $\phi_B$  will be depth dependent. The 20 experimental data for the parameter determination consist of well-log measurements from a brine containing zone, laboratory Formation Resistivity Factor (FRF) measurements and brine saturation experiments. The log information from the brine containing zone is used to 25 correlate the local parameter  $\phi_C$  to suitable logs/log combinations, as is known to those skilled in the art of well logging.  $\sigma_C,~\lambda_B$  and  $\varphi_B$  and the correlation of  $\phi_C$  to suitable logs/log combinations can be used in hydrocarbon bearing formations. From the well-log, the above 30 relationship and the indicated parameters, the volume

faction of brine and thus also the volume fraction of hydrocarbon is determined as a function of depth.

	petro	petrophysical	data	ц.	parameters	,,	incoherence	rence	incohe	incoherence
			,				for present	esent	for prior	ior art
sample	· •	. K [mD]	Ô	ø	o C	h <sub>B</sub>	invention $\sum \Delta_{c_o-c_o}^i$	tion $\sum \Delta_{\psi}^2$	$\sum \Delta^{l}_{c_o-c_{\bullet}}$	$\sum_{\mathbf{v}_{\mathbf{v}_{\mathbf{v}}}} \Delta_{\mathbf{v}_{\mathbf{v}}}^2$
No.			[eq/1]	-	[mS/cm]					
1	0.239	629	0.017	0.0532	2.9581	0.2016	0.002	0.018	0.009	0.026
8	0.212	105	0.052	0.1822	1.3537	0.2038	0.002	0.035	0.061	0.099
м	0.231	397	0.052	0.1376	1.8607	0.2517	0.002	0.014	0.031	0.097
4	0.080	1.34	0.26	0.1323	1.3325	0.1788	0.010	0.119	0.045	0.010
Ŋ	0.154	55	0.2	0.3216	1.4851	0.3752	0.005	0.042	0.149	0.128
ø	0.215	29	0.095	0.3467	1.0193	0.1370	0.003	0.026	0.206	0.257
7	0.171	3.5	0.053	0.3779	0.7847	0.1200	0.008	0.016	0.312	0.446
©	0.171	7.66	0.053	0.3119	0.8994	0.1274	0.007	0.018	0.234	0.366
o.	0.199	57	0.085	0.3935	1.0627	0.1613	0.007	0.025	0.276	0.351
10	0.125	0.042	0.253	0.432	0.3113	0.0255	0.029	0.074	0.483	0.262
11	0.125	0.0106	0.253	0.4007	0.2590	0.0233	0.037	0.057	0.348	0.203
12	0.110	1.86	0.28	0.5855	0.8601	0.1326	0.018	0.010	0.760	0.542
13	0.110	0.3	0.28	0.5998	1.0802	0.1286	0.020	0.016	0.868	0.589
14	0.110	2.08	0.28	0.5815	1.4104	0.1998	0.054	0.029	0.854	0.539
15	0.092	0.128	0.41	0.509	0.6025	0.0402	0.027	0.025	0.917	0.449
16	0.103	0.024	0.67	0.7202	1.4537	0.1103	0.010	0.013	1.263	0.529
17	0.140	0.575	0.33	0.7763	1.5783	.7760.0	0.050	0.013	1.606	0.919
18	0.259	3.78	0.59	0.7033	2.8964	0.1357	0.012	0.007	1.173	0.507
19	0.259	17.1	0.59	0.6408	2.8227	0.1484	0.010	0.012	0.823	0.376

	petro	petrophysical	data	ц	parameters	10	incohe	incoherence	incoherence for	ance for
							for p	for present	prior art	art
sample	æ	K [mD]	6	ø	p	h	invention $\sum \Delta_{c_o-c_o}^2 \sum \Delta_{\psi}^2$		$\sum \Delta^{i}_{c_o \cdot c_o}$	$\sum \Delta_{\psi}^2$
No.			[eq/1]		[mS/cm]					:
20	0.259	44.8	0.59	0.5978	3.4771	0.2113	0.003	0.023	0.674	0.286
21	0.238	315	0.29	0.6984	1.3859	0.1812	0.009	0.031	0.632	0.546
22	0.225	1.92	0.72	0.7651	2.4172	0.0766	0.016	0.013	1.808	0.637
23	0.242	54.3	1.04	0.7306	4.0758	0.1165	0.022	0.011	1.640	0.490
24	0.216	0.546	0.81	0.7751	2.6418	0.0659	0.021	0.014	2.103	0.703
25	0.187	0.0348	1.27	0.7995	3,4510	0.0726	0.005	0.040	0.490	0.369
26	0.229	1.53	1.47	0.7491	5.0425	0.0887	0.042	0.026	2.155	0.470
27	0.209	0.263	1.48	0.7656	5.1455	0.0884	0.0884 0.001 0.067 0.495	0.067	0.495	0.326

TABLE

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and

#### CLAIMS

- 1. A method of determining a parameter selected from the electrical conductivity and the volume fraction of a component in a composition comprising a plurality of components, the method comprising:
- 5 measuring the electrical conductivity of the composition;
  - selecting a relationship between the electrical conductivity of the composition and a plurality of composition parameters including, for each component, physical parameters representing the electrical conductivity and the volume fraction of the component, the components being substantially equally represented in

said relationship by means of said physical parameters;

- determining said selected parameter from said relationship and the measured conductivity of the composition.
  - 2. The method of claim 1, wherein said plurality of composition parameters includes at least one fitting parameter, and wherein each fitting parameter is determined by applying said relationship to a data set obtained by measuring the electrical conductivity of at least one sample representative for said composition for various magnitudes of at least one of said parameters.
- 3. The method of claim 1 or 2, wherein said plurality of parameters includes an auxiliary parameter depending on the geometrical configuration of the components in the composition.
- The method of claim 3, wherein said relationship is
   selected to be

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$$(\sigma_{\text{eff}} - \sigma_0) \cdot (L\sigma_{\text{eff}} + (1-L)\sigma_0)^{-1} = \sum \phi_k (\sigma_k - \sigma_0) \cdot (L\sigma_k + (1-L)\sigma_0)^{-1}$$

wherein  $\sigma_0$  represents the auxiliary parameter in the form of a conductivity tensor

k = 1 ... N, N being the number components  $\sigma_{\mbox{\scriptsize eff}}$  represents the conductivity tensor of the sample

 $\boldsymbol{\sigma}_k$  represents the conductivity tensor of component k

 $\phi_k$  represents the volume fraction of component k L represents a depolarisation tensor

- 5. The method of claim 3 or 4, wherein said auxiliary parameter is selected so as to be a function of a plurality of variables, each variable depending on the conductivity of one of said components and a mixing coefficient, whereby the mixing coefficients depend on the geometrical configuration of the components in the composition.
  - 6. The method of any of claims 3-5, wherein said auxiliary parameter is selected to be

 $\sigma_0 = \sum h_k \sigma_k$ 

wherein  $\sigma_0$  represents the auxiliary parameter in the form of a conductivity tensor representative for the conductivity in the three principal directions;

k=1 ... N, N being the number of components;  $\sigma_k$  represents the conductivity tensor of component k:

 $\mathbf{h}_{k}$  represents the mixing coefficient tensor pertaining to component  $\mathbf{k}$ ;

- 7. The method of claim 5 or 6, wherein said mixing coefficients are selected so that the sum of the mixing coefficients substantially equals unity.
- 8. The method of any of claims 5-7, wherein said mixing coefficients are non-negative.

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- 9. The method of any of claims 5-8, wherein each mixing coefficient is selected to be a function of at least the volume fraction of the component pertaining to said mixing coefficient.
- 5 10. The method of claim 9, wherein said function is a monotonous increasing function in the volume fraction of the component pertaining to the mixing coefficient.
  - 11. The method of claim 9 or 10, wherein said function is selected so that the mixing coefficient vanishes for vanishing volume fraction of the component pertaining to the mixing coefficient.
    - 12. The method of any of claims 5-11, wherein each mixing coefficient is selected as

 $h_k = \lambda_k \phi_k V_k \quad (\sum \lambda_n \phi_n V_n)^{-1}$ 

wherein  $k,n = 1 \dots N$ , N being the number components in said plurality of components

 $\boldsymbol{\lambda}_k$  represents a percolation rate tensor pertaining to component k

- $\varphi_k$  represents the volume fraction of component k  $\nu_{k,\,n}$  represents a percolation exponent pertaining to component k,n
  - 13. The method of claim 12, wherein at least one of  $h_k$ ,  $\lambda$  and  $\nu$  forms a fitting parameter.
- 14. The method of any of claims 4-13, wherein the depolarisation tensor is positive.
  - 15. The method of any of claims 4-14, wherein the depolarisation tensor has unit trace.
  - 16. The method of any of claims 4-15, wherein the depolarisation tensor equals 1/3 times the unit tensor.
- 30 17. The method of any of claims 1-16, wherein the step of determining each fitting parameter by applying said relationship to the data set is carried out through an iterative process.

- 18. The method of claim 17, wherein the iterative process includes repeatedly applying said relationship in a minimisation scheme.
- 19. The method of claim 18, wherein the minimisation scheme is applied to a mismatch between the measured electrical conductivities of said components and the electrical conductivities of the components as determined through said relationship.
- 20. The method of any of claims 1-19, wherein said composition includes an earth formation.
  - 21. The method of claim 20, wherein said earth formation includes at least one of rock, brine, hydrocarbon fluid and clay.
- 22. The method of claim 21, wherein said parameter which is determined forms the volume fraction of one of the hydrocarbon fluid and the brine.

## INTERNATIONAL SEARCH REPORT

Inte onal Application No PCT/EP 96/01234

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A. CLASSIE IPC 6	FICATION OF SUBJECT MATTER G01V3/38 G01N33/24		
According to	International Patent Classification (IPC) or to both national class	sification and IPC	
	SEARCHED		
Minimum do IPC 6	cumentation searched (classification system followed by classific GO1V GO1N	ation symbols)	
Documentation	on searched other than minimum documentation to the extent tha	t such documents are included in the fields sea	rched
Electronic da	ta base consulted during the international search (name of data b	ase and, where practical, search terms used)	
C. DOCUMI	ENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the	relevant passages	Relevant to claim No.
Α	US,A,4 398 151 (VINEGAR ET AL.) 1983 see the whole document	9 August	1,20
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X Further	er documents are listed in the continuation of box C.	X Patent family members are listed in	annex.
"A" document consider artist of filing da "L" document which is citation "O" document other m	nt which may throw doubts on priority claim(s) or streed to establish the publication date of another or other special reason (as specified) nt referring to an oral disclosure, use, exhibition or	"T" later document published after the inter or priority date and not in conflict with cited to understand the principle or the invention.  "X" document of particular relevance; the cannot be considered novel or cannot involve an inventive step when the document of particular relevance; the cannot be considered to involve an involve an involve an invention to combined with one or moments, such combination being obvious in the art.  "&" document member of the same patent of	n the application but ony underlying the laimed invention be considered to urment is taken alone laimed invention entive step when the re other such docu- s to a person skilled
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	ailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Russwijk Td. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer  Bosma, R	

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# INTERNATIONAL SEARCH REPORT

Inte onal Application No
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